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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/521,044	. 12/07/2005	Walid Nagib Aboul-Hosn	OMA003-US1	6603
Jonathan Spangler 2875 Kalmia Place San Diego, CA 92104			EXAMINER	
			FLORY, CHRISTOPHER A	
			ART UNIT	PAPER NUMBER
	•		3762	
SHORTENED STATUTORY PERIOD OF RESPONSE		MAIL DATE	DELIVERY MODE	
3 MONTHS		01/25/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

If NO period for reply is specified above, the maximum statutory period will apply and will expire 6 MONTHS from the mailing date of this communication.

	Application No.	Applicant(s)			
	10/521,044	ABOUL-HOSN, WALID NAGIB			
Office Action Summary	Examiner	Art Unit			
	Christopher A. Flory	3762			
The MAILING DATE of this communication ap Period for Reply	pears on the cover sheet with t	the correspondence address			
A SHORTENED STATUTORY PERIOD FOR REPLEWHICHEVER IS LONGER, FROM THE MAILING ID. - Extensions of time may be available under the provisions of 37 CFR 1. after SIX (6) MONTHS from the mailing date of this communication. - If NO period for reply is specified above, the maximum statutory period. - Failure to reply within the set or extended period for reply will, by statut Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNICA .136(a). In no event, however, may a reply d will apply and will expire SIX (6) MONTHS te, cause the application to become ABAND	TION. be timely filed from the mailing date of this communication. DONED (35 U.S.C. § 133).			
Status		•			
1)⊠ Responsive to communication(s) filed on 11 .	January 2005.				
	is action is non-final.	•			
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closed in accordance with the practice under					
Disposition of Claims					
4)⊠ Claim(s) <u>1-15</u> is/are pending in the application	n. '				
4a) Of the above claim(s) is/are withdra					
5) Claim(s) is/are allowed.					
6)⊠ Claim(s) <u>1-15</u> is/are rejected.					
7) Claim(s) is/are objected to.					
8) Claim(s) are subject to restriction and/	or election requirement.				
Application Papers					
9) The specification is objected to by the Examin	er.				
10) The drawing(s) filed on is/are: a) ac		the Examiner.			
Applicant may not request that any objection to the					
Replacement drawing sheet(s) including the correct					
11)☐ The oath or declaration is objected to by the E	xaminer. Note the attached O	ffice Action or form PTO-152.			
Priority under 35 U.S.C. § 119					
12)⊠ Acknowledgment is made of a claim for foreign a)⊠ All b)□ Some * c)□ None of:	n priority under 35 U.S.C. § 11	9(a)-(d) or (f).			
1. Certified copies of the priority documents have been received.					
2. Certified copies of the priority documents have been received in Application No					
3. Copies of the certified copies of the price	ority documents have been red	ceived in this National Stage			
application from the International Burea	au (PCT Rule 17.2(a)).				
* See the attached detailed Office action for a lis	t of the certified copies not rec	eived.			
·					
Attachment(s)					
1) Notice of References Cited (PTO-892)	4) Interview Sumi				
2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08)		ail Date mal Patent Application			
Paper No(s)/Mail Date	6) Other:				

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DETAILED ACTION

Claim Rejections - 35 USC § 102

1. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

- (b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.
- (e) the invention was described in a patent granted on an application for patent by another filed in the United States before the invention thereof by the applicant for patent, or on an international application by another who has fulfilled the requirements of paragraphs (1), (2), and (4) of section 371(c) of this title before the invention thereof by the applicant for patent.

The changes made to 35 U.S.C. 102(e) by the American Inventors Protection Act of 1999 (AIPA) and the Intellectual Property and High Technology Technical Amendments Act of 2002 do not apply when the reference is a U.S. patent resulting directly or indirectly from an international application filed before November 29, 2000. Therefore, the prior art date of the reference is determined under 35 U.S.C. 102(e) prior to the amendment by the AIPA (pre-AIPA 35 U.S.C. 102(e)).

2. Claims 1-3, 8 and 12 are rejected under 35 U.S.C. 102(b) as being anticipated by Izraelev (US Patent 5,685700, hereinafter Izraelev'700).

Regarding claims 1-3, Izraelev'700 discloses a blood pump for percutaneous introduction into a patient (TITLE) comprising a pump housing (Fig. 2, housing 11) having at least two blood inlets disposed on opposing ends of said pump housing (ABSTRACT; column 2, lines 13-16; Fig. 2 inlets 16 and 17); at least one blood outlet pump disposed between the blood inlets (ABSTRACT; Fig. 3, outlets 18 and 19); a rotor chamber

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extending between the inlets and outlets (Fig. 2, chamber 12); and a rotor operable to draw blood into the inlets and direct blood to said outlets (ABSTRACT; Fig. 2, rotor 20).

Regarding claims 8 and 12, Izraelev'700 discloses cooperating magnets to position said rotor within said pump housing and a control system to control said motor (abstract; column 1, lines 22-25; column 4, lines 37-56).

3. Claims 1-10 and 12-15 are rejected under 35 U.S.C. 102(e) as being anticipated by Schulte Eistrup et al. (US Patent 6,752,602, hereinafter Eistrup'602).

Regarding claims 1-3, Eistrup'602 discloses a blood pump for percutaneous introduction into a patient (TITLE) comprising a pump housing (Fig. 1, housing 10) having at least two blood inlets disposed on opposing ends of said pump housing (ABSTRACT; Fig. 1, inlets 14 and 16); at least one blood outlet pump disposed between the blood inlets (ABSTRACT; Fig. 1, outlet 18); a rotor chamber extending between the inlets and outlets (Fig. 1, chamber 12); and a rotor operable to draw blood into the inlets and direct blood to said outlets (ABSTRACT; Fig. 1, rotor 20).

Regarding claim 4, Eistrup'602 discloses an outflow cannula coupled to said blood outlet (column 8, line 67).

Regarding claim 9, Eistrup'602 discloses thrust bearings (abstract; column 3, lines 10-13; Fig. 1, bearing pair 34).

Regarding claims 8, 10 and 12, Eistrup'602 discloses cooperating magnets to position said rotor within said pump housing and a control system to control said motor (column 3, lines 35-42; column 4, lines 45-47; Fig. 1, rotor magnets 28).

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Regarding claims 13-15, Eistrup'602 discloses a rechargeable, subcutaneous battery (column 9, lines 5-10).

Claim Rejections - 35 USC § 103

- 4. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:
 - (a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negatived by the manner in which the invention was made.
- 5. Claim 11 is rejected under 35 U.S.C. 103(a) as being unpatentable over Izraelev'700 in view of Viole et al. (US Patent 6,610,004, hereinafter Viole'004).

Regarding claim 11, Izraelev'700 discloses the invention substantially as claimed but does not expressly disclose that the blood pump further include a protective cage disposed over at least one of said blood inlets. In the same field of endeavor, Viole'004 teaches a housing with openings having a cage-like arrangement to shield the pump blades from damaging endothelial lining (column 8, lines 23-30). Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to modify the system of Izraelev'700 with the cage structure of Viole'004 to provide Izraelev'700 with the same advantages of preventing damage to the endothelial lining.

6. Claims 13-15 are rejected under 35 U.S.C. 103(a) as being unpatentable over lzraelev'700.

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Izraelev'700 discloses the invention substantially as claimed, but does not expressly disclose a subcutaneous rechargeable battery. However, it is well known in the art to use subcutaneous rechargeable batteries to power implanted devices so that the device may be used beyond a single battery lifetime without need for explant or additional surgery which might further injure the patient or damage the implanted device. Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to modify the system as taught by Izraelev'700 with a rechargeable subcutaneous battery to provide the advantage of being able to extend the usage of the implanted device without the need to perform surgery and explant the device, causing further damage to the device and the patient, as is well known in the art.

7. Claims 5 and 7 are rejected under 35 U.S.C. 102(e) as anticipated by Eistrup'602 or, in the alternative, under 35 U.S.C. 103(a) as obvious over Eistrup'602 in view of Viole'004 or in view of Jarvik (US Patent 4,994,078, hereinafter Jarvik'078).

Regarding claims 5 and 6, the cannula disclosed in Eistrup'602 is capable of extending across a valve or through an opening created in the aorta. Alternatively, in the same field of endeavor, Viole'004 teaches a cannula passing through the aorta so that blood may be directed from a first to a second blood vessel (column 16, lines 31-56). Similarly, Jarvik'078 teaches inserting a cannula into the aortic wall and across the aortic valve to make the valve incompetent and permit leaks in a suture line to be detected and repaired before connection of the intraventricular blood pump is made (column 14, lines 34-41). Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to modify the system of Eistrup'602 with the

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cannula placed across a valve or the aorta to provide the Eistrup'602 system with the same advantages of directing blood across blood vessels or detecting and repairing suture leaks before a blood pump is connected permanently.

8. Claim 6 is rejected under 35 U.S.C. 102(e) as anticipated by Eistrup'602 or, in the alternative, under 35 U.S.C. 103(a) as obvious over Eistrup'602 in view of Zafirelis et al. (US Patent 6,808,508, hereinafter Zafirelis'508).

Regarding claim 6, Eistrup'602 discloses a cannula capable of being placed through an opening in the atrial septum. Alternatively, Zafirelis'508 teaches a transseptal cannula placed through the atrial septum for returning oxygenated blood to the arterial system of the patient (ABSTRACT). Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to modify the system of Eistrup'602 with the transseptal cannula as taught by Zafirelis'508 in order to provide the Eistrup'602 system with the same advantage of returning oxygenated blood to the arterial system of a patient (motivation to combine provided by Zafirelis'508 ABSTRACT).

9. Claim 11 is rejected under 35 U.S.C. 103(a) as being unpatentable over Eistrup'602 in view of Viole'004 et al.

Regarding claim 11, Eistrup'602 discloses the invention substantially as claimed but does not expressly disclose that the blood pump further include a protective cage disposed over at least one of said blood inlets. In the same field of endeavor, Viole'004 teaches a housing with openings having a cage-like arrangement to shield the pump blades from damaging endothelial lining (column 8, lines 23-30). Therefore, it would have been obvious to one of ordinary skill in the art at the time of the invention to modify

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the system of Eistrup'602 with the cage structure of Viole'004 to provide Izraelev'700

with the same advantages of preventing damage to the endothelial lining.

Conclusion

Any inquiry concerning this communication or earlier communications from the

examiner should be directed to Christopher A. Flory whose telephone number is (571)

272-6820. The examiner can normally be reached on M - F 8:30 a.m. to 5:00 p.m..

If attempts to reach the examiner by telephone are unsuccessful, the examiner's

supervisor, Angela Sykes can be reached on (571) 272-4955. The fax phone number

for the organization where this application or proceeding is assigned is 571-273-8300.

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Christopher A. Flory

20 December 2006

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